

Title (en)  
GENE DISRUPTION METHODOLOGIES FOR DRUG TARGET DISCOVERY

Title (de)  
VERFAHREN ZUR GENUNTERBRECHUNG ZUM ERKENNEN VON ANGRIFFSZIELEN VON MEDIKAMENTEN

Title (fr)  
PROCEDES DE DISRUPTION DE GENES POUR IDENTIFICATION DE CIBLE DE MEDICAMENT

Publication  
**EP 1348027 A2 20031001 (EN)**

Application  
**EP 01991419 A 20011226**

Priority  
• US 0149486 W 20011226  
• US 25912800 P 20001229  
• US 79202401 A 20010220  
• US 31405001 P 20010822

Abstract (en)  
[origin: WO02053728A2] The present invention provides methods and compositions that enable the experimental determination as to whether any gene in the genome of a diploid pathogenic organism is essential, and whether it is required for virulence or pathogenicity. The methods involve the construction of genetic mutants in which one allele of a specific gene is inactivated while the other allele of the gene is placed under conditional expression. The identification of essential genes and those genes critical to the development of virulent infections, provides a basis for the development of screens for new drugs against such pathogenic organisms. The present invention further provides Candida albicans genes that are demonstrated to be essential and are potential targets for drug screening. The nucleotide sequence of the target genes can be used for various drug discovery purposes, such as expression of the recombinant protein, hybridization assay and construction of nucleic acid arrays. The uses of proteins encoded by the essential genes, and genetically engineered cells comprising modified alleles of essential genes in various screening methods are also encompassed by the invention.

IPC 1-7  
**C12N 15/10**

IPC 8 full level  
**G01N 33/50** (2006.01); **A61K 45/00** (2006.01); **A61K 48/00** (2006.01); **A61P 31/10** (2006.01); **C07K 14/37** (2006.01); **C07K 14/40** (2006.01); **C07K 16/14** (2006.01); **C12N 1/15** (2006.01); **C12N 1/19** (2006.01); **C12N 1/21** (2006.01); **C12N 5/10** (2006.01); **C12N 15/09** (2006.01); **C12N 15/10** (2006.01); **C12P 21/02** (2006.01); **C12P 21/08** (2006.01); **C12Q 1/02** (2006.01); **C12Q 1/68** (2006.01); **G01N 33/15** (2006.01); **G01N 33/53** (2006.01); **G01N 37/00** (2006.01)

CPC (source: EP US)  
**A61P 31/10** (2017.12 - EP); **C07K 14/37** (2013.01 - EP US); **C07K 14/40** (2013.01 - EP US); **C12N 1/145** (2021.05 - EP US); **C12N 1/165** (2021.05 - EP US); **C12N 15/1082** (2013.01 - EP US); **C12R 2001/645** (2021.05 - EP US); **C12R 2001/725** (2021.05 - EP US)

Citation (search report)  
See references of WO 02053728A2

Cited by  
CN108823234A

Designated contracting state (EPC)  
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE TR

DOCDB simple family (publication)  
**WO 02053728 A2 20020711**; **WO 02053728 A3 20030320**; **WO 02053728 A9 20030530**; CA 2432902 A1 20020711; EP 1348027 A2 20031001; JP 2005514899 A 20050526; US 2003180953 A1 20030925

DOCDB simple family (application)  
**US 0149486 W 20011226**; CA 2432902 A 20011226; EP 01991419 A 20011226; JP 2002555238 A 20011226; US 3258501 A 20011220